

A new neutron-gamma density measurement method using mass attenuation coefficient function*

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While traditional gamma-gamma density (GGD) logging technology is widely utilized, its potential environmental risks have prompted the development of more environmentally friendly neutron-gamma density (NGD) logging technology. However, NGD measurements are influenced by both neutron and gamma radiation. In the logging environment, variations in formation composition indicate different elemental compositions, which affect neutron-gamma reaction cross-sections and gamma generation. Compared to traditional gamma sources such as Cs-137, these changes can significantly impact the generation and transport of neutron-induced inelastic gamma rays, posing challenges for accurate measurements. To address this, a novel method is proposed, the method incorporates the mass attenuation coefficient function to account for the effects of various lithologies and pore contents on gamma-ray attenuation and therefore can achieve more accurate density measurements by clarifying the transport processes of inelastic gamma rays with varying energy and spatial distributions in varied logging environments. The method avoids the complex correction of the neutron transport and is verified through Monte Carlo simulations for its applicability across various lithologies and pore contents, showing that the absolute density errors are less than 0.02 g/cm³ in clean formations and demonstrating good accuracy. The research not only clarifies the NGD mechanism but also provides theoretical guidance for the application of NGD logging methods. Further research will be conducted regarding extreme environmental conditions and tool calibration.

Keywords: Neutron-gamma density, Mass attenuation coefficient, Monte Carlo simulation

I. INTRODUCTION

In field of petroleum exploration, traditional gamma-gamma density (GGD) logging technology has played a crucial role for many years [1–5]. However, with growing awareness of environmental protection, the risks of pollution and operational safety associated with GGD technology have become increasingly apparent, posing challenges to its further development. In this context, neutron-gamma density (NGD) logging technology has emerged as new research focus due to its advantage of environmental protection and controllability [6–10]. GGD relies on the transport of monoenergetic gamma rays from the source to the detectors, while NGD is based on the transport of neutron-induced gamma rays, whose energy exhibits uncertainty. In NGD, the gamma rays detected by the detector are influenced by neutron transport from the neutron source to the point of the gamma ray producing neutron interaction in the formation and by the subsequent transport of the gamma rays from their source to the gamma-ray detector. To eliminate the influence of thermal neutron effects, density measurements are conducted using inelastic gamma rays produced by high-energy neutrons [11, 12]. Once neutrons are emitted, they undergo inelastic scattering reactions with isotopes of crucial elements in the medium in a few microseconds, producing inelastic gamma rays. These gamma rays are less influenced by neutron transport, enabling them to more accurately reflect formation characteristics, making them more suitable for density measurements. However, evaluating inelastic gamma rays can be tricky because it depends

on both neutron and gamma physics and undergoes multiple physics processes simultaneously. Hence in this case, the entanglement between neutron and gamma transport increases the complexity of the measurement and its sensitivity to its environment. The generation and attenuation of inelastic gamma rays are directly influenced by environment factors such as lithology, porosity, and fluid properties. It is critical to understand both neutron and gamma physics pertaining to changing environments in order to develop an accurate method for NGD technology [13–15].

The development of neutron-gamma density (NGD) technology has been actively ongoing for the past few decades. For instance, Odom et al. used inelastic gamma rays for density measurements, which advanced density logging technology based on the neutron-gamma coupled field theory [16, 17]. However, this method is affected by neutron transport, and neutron transport correction needs to be considered in subsequent studies. Jacobson et al. developed a correction technique that employs capture gamma count ratio to obtain a compensated inelastic gamma ratio, achieving the density measurements [18]. Zhang et al. developed a density method by using the inelastic gamma-count ratio and the fast-neutron count to avoid neutron correction [19]. Luyckx et al. approximated the initial inelastic gamma flux by fast neutron counts for density measurements [20]. Wang et al. created a correction model utilizing epithermal neutrons, and divided the inelastic gamma-ray energy spectrum into high- and low-energy windows to reduce the influence of pair production [21]. Additionally, Zhang et al. introduced an adaptive method for obtaining inelastic gamma spectra while environment changes and integrated capture correction for density measurement [22]. While these studies have made progress, most researchers focused more on analyzing the neutron transport process and less on the dynamic changes in gamma attenuation process. Inelastic gamma rays generated by neutron-induced reactions exist in formations in a non-

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65 monoenergetic distribution, whereas chemical sources like
 66 Cs-137 generate monoenergetic gamma in a homogeneous
 67 manner. Furthermore, typical neutron-induced gamma rays
 68 can reach energies up to 8 MeV [23]. Pair production needs
 69 to be considered as it plays a vital role in the neutron-induced
 70 gamma transport process. These factors contribute to the
 71 complexity of gamma attenuation. Previous neutron-gamma
 72 density (NGD) and gamma-gamma density (GGD) provide
 73 possibilities for density measurements. However, most of
 74 previous NGD methods consider mass attenuation coefficient
 75 as a constant, this limits the accuracy because it is closely re-
 76 lated to the formation composition. In our work, we introduce
 77 mass attenuation coefficient as a function related to formation
 78 lithologies and pore contents, in order to accurately depict
 79 the intricate interaction mechanisms between radiation and
 80 formation, which is essential in obtaining accurate formation
 81 density. This can also provide a new approach to complement
 82 previous methods.

83 The manuscript is organized as follows: Section 2 in-
 84 troduces the method and presents the development process.
 85 Also, a pulsed neutron density tool is described, which later
 86 is employed for concept verification. Section 3 presents the
 87 results from different simulated scenarios, demonstrating the
 88 method's effectiveness. Finally, Conclusions are drawn in
 89 Section 4.

II. METHOD

90 The development of the method is shown in Fig.1: Box1
 91 reviews the coupled field theory of neutron-gamma density
 92 (NGD) measurement, which is the foundation for the pro-
 93 posed method as it depicts inelastic gamma rays' distribution.
 94 Box2 is key to the method: a function for mass attenuation
 95 coefficient is developed, which is then used to derive den-
 96 sity. Certain key parameters, such as the hydrogen index can-
 97 not be directly expressed in this mathematical form, instead,
 98 they are obtained through tool measurement. Box3 presents
 99 analysis of physical parameters using a real NGD tool under
 100 development stage. Extensive Monte Carlo simulations are
 101 conducted to establish a quantitative relationship between de-
 102 tector responses and formation physical parameters, which is
 103 consequently utilized to obtain these key parameters. Finally,
 104 density is calculated. Overview of the method is illustrated in
 105 Fig.1.

A. Coupled field theory of NGD Measurement

109 NGD logging technology relies on inelastic gamma rays to
 110 measure formation density [24–29]. The distribution of in-
 111 elastic gamma rays involves two interconnected links of neu-
 112 tron and gamma transport [30–32]. Detailed process is dis-
 113 cussed as below:

114 The pulsed neutron source emits 14MeV fast neutrons. In
 115 a spherical model, according to neutron diffusion theory, the
 116 distribution of fast neutrons can be described as follows:

$$117 \quad \Phi_n = \frac{Q}{4\pi D_n r} \exp\left(-\frac{r}{L_n}\right) \quad (1)$$

118 where Q is the number of neutrons emitted by the neutron
 119 source per second, D_n is the neutron diffusion coefficient, L_n
 120 is the fast neutron deceleration length, and r is the distance
 121 between the pulsed neutron source and the neutron detector.
 122 The inelastic gamma rays recorded by a detector with a radius
 123 R , can be described by the following equation [21]:

$$124 \quad \Phi_{in}(R) = \frac{i\Sigma_{in} Q \int_0^\infty \exp\left(-\frac{r}{L_n}\right) \exp(-\rho\mu_m |r - R|) dr}{4\pi D_n R} \quad (2)$$

125 where i is the average number of inelastic gamma rays after
 126 neutron enters the formation, Σ_{in} is the inelastic scattering
 127 cross section, ρ is the formation density, and μ_m is the total
 128 mass attenuation coefficient. The inelastic gamma rays within
 129 the distance from the source R are recorded [33], and Eq.(2)
 130 can be written as:

$$131 \quad \Phi_{in}(R) = \frac{i\Sigma_{in} Q}{4\pi D_n R} \frac{\exp\left(-\frac{R}{L_n}\right) - \exp(-\rho\mu_m R)}{\rho\mu_m - \frac{1}{L_n}} \quad (3)$$

132 According to the Lagrange interpolation method, Eq.(3)
 133 can be simplified as follows:

$$134 \quad \Phi_{in}(R) = \frac{i\Sigma_{in} Q}{4\pi D_n} \exp(-R\xi) \quad (4)$$

135 where ξ belongs to $(\rho\mu_m, \frac{1}{L_n})$, which can be written as: $\xi =$
 136 $\frac{1}{L_n} - \alpha(\frac{1}{L_n} - \rho\mu_m)$, $\alpha \in (0, 1)$.

137 Assuming that the source distances of the near and far
 138 gamma detectors are L_1 and L_2 ($L_1 < L_2$), the following
 139 equations can be obtained:

$$140 \quad \begin{cases} \Phi_{in}(L_1) = \frac{i\Sigma_{in} Q}{4\pi D_n} \exp(-L_1\xi_1) \\ \Phi_{in}(L_2) = \frac{i\Sigma_{in} Q}{4\pi D_n} \exp(-L_2\xi_2) \end{cases} \quad (5)$$

141 where:

$$142 \quad \begin{cases} \xi_1 = \frac{1}{L_n} - \alpha_1(\frac{1}{L_n} - \rho\mu_m), \alpha_1 \in (0, 1) \\ \xi_2 = \frac{1}{L_n} - \alpha_2(\frac{1}{L_n} - \rho\mu_m), \alpha_2 \in (0, 1) \end{cases} \quad (6)$$

143 The logarithm of ratio of near and far inelastic gamma
 144 counts is as follows:

$$145 \quad \begin{aligned} \ln(RIN) &= \ln\left(\frac{\Phi_{in}(L_1)}{\Phi_{in}(L_2)}\right) \\ &= \frac{L_1\alpha_1 - L_2\alpha_2 + L_2 - L_1}{L_n} + (L_2\alpha_2 - L_1\alpha_1)\rho\mu_m \end{aligned} \quad (7)$$

146 Suppose $a = L_1\alpha_1 - L_2\alpha_2 + L_2 - L_1$, $b = L_2\alpha_2 -$
 147 $L_1\alpha_1$, Eq.(7) can be reorganized:

$$148 \quad \ln(RIN) = \frac{a}{L_n} + b\rho\mu_m \quad (8)$$

149 Eq. (8) shows that the response of gamma-ray detector is
 150 related to the L_n , ρ , μ_m , and the value of a , b . The fast neu-

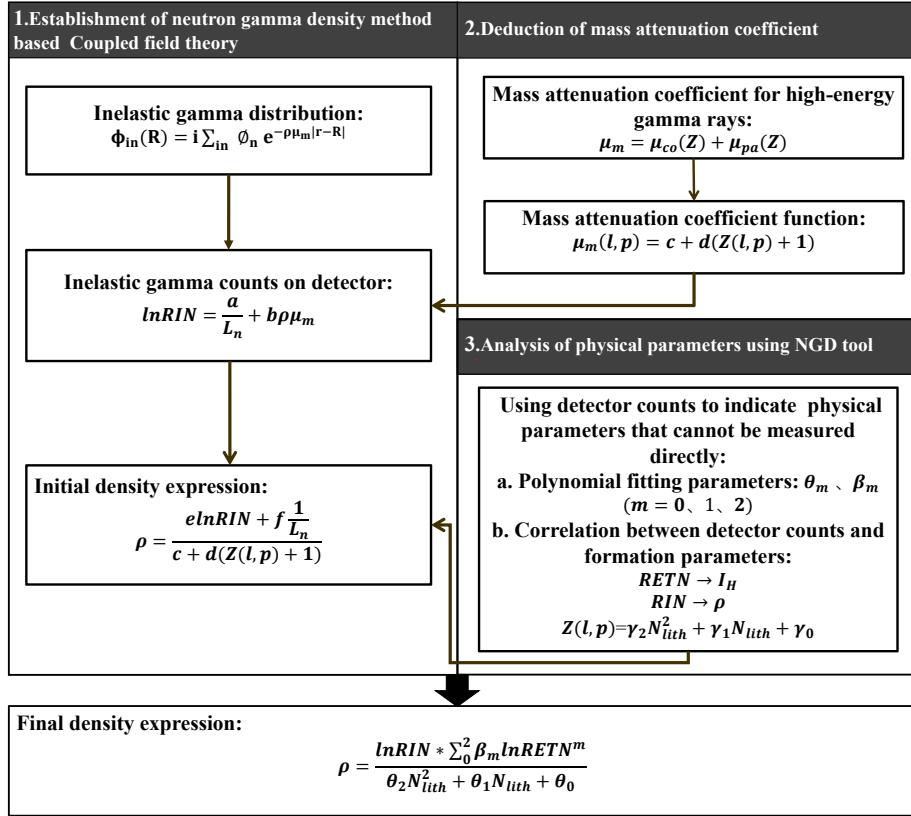


Fig. 1. (Color online) Overview of the proposed method

151 tron deceleration length, L_n in typical formations can be char-
152 acterized by the formation density ρ , the hydrogen index I_H ,
153 and the initial energy of fast neutron E_0 [34]:

$$154 \quad L_n = k \sqrt{\frac{1}{I_H + c}} \sqrt{\frac{1}{\rho}} \sqrt{\ln(E_0)} \quad (9)$$

155 The key parameter in Eq.(8) is the mass attenuation
156 coefficient μ_m , which directly reflects the attenuation of
157 gamma rays in formations. The section below will focus on
158 analyzing this parameter.

B. NGD method development

160 The attenuation of gamma rays in formations is closely linked
161 to formation density and the mass attenuation coefficient.
162 This attenuation process is primarily influenced by various
163 physics processes, including the Photoelectric effect, Com-
164 ton effect, and Pair production. Especially for high-energy
165 gamma rays, Compton effect and Pair production are the main
166 factors affecting their attenuation. Thus, the total attenuation
167 coefficient μ_m can be expressed as follows:

$$168 \quad \mu_m = \mu_{co} + \mu_{pa} \quad (10)$$

169 where μ_{co} is the mass attenuation coefficient for Compton

170 effect, μ_{pa} is the mass attenuation coefficient for pair produc-
171 tion.

172 According to the principles of Compton effect and Pair pro-
173 duction [35, 36], the mass attenuation coefficient of Compton
174 effect and Pair production can be expressed as:

$$175 \quad \mu_{co} = 2\pi(r_0)^2 \left\{ \frac{1+\eta}{\eta^2} \left[\frac{2(1+\eta)}{1+2\eta} - \frac{1}{\eta} \ln(1+2\eta) \right] \right. \\ \left. + \frac{1}{2\eta} \ln(1+2\eta) - \frac{1+3\eta}{(1+2\eta)^2} \right\} N_A \frac{Z}{A} \quad (11)$$

$$176 \quad \mu_{pa} = \frac{(r_0)^2 N_A}{137} \left(\frac{28}{9} \ln(2\eta) - \frac{218}{27} \right) (Z+1) \frac{Z}{A} \quad (12)$$

177 where N_A is Avogadro constant, r_0 is the classical electron
178 radius ($r_0=2.818*10^{-13}\text{cm}$), $\eta = \frac{E_\gamma}{m_e c^2}$, E_γ is the gamma-
179 ray energy, m_e is the electron rest mass ($m_e=9.110*10^{-31}\text{kg}$),
180 c is the speed of light in a vacuum ($c=2.998*10^8\text{m/s}$), Z is
181 the atomic number, A is the atomic weight. The total mass
182 attenuation coefficient can be rewritten as:

$$183 \quad \mu_m = \pi(r_0)^2 \left\{ \frac{1+\eta}{\eta^2} \left[\frac{2(1+\eta)}{1+2\eta} - \frac{1}{\eta} \ln(1+2\eta) \right] \right. \\ \left. + \frac{1}{2\eta} \ln(1+2\eta) - \frac{1+3\eta}{(1+2\eta)^2} \right\} N_A \\ + \frac{(r_0)^2 N_A}{274} \left(\frac{28}{9} \ln(2\eta) - \frac{218}{27} \right) (Z+1) \quad (13)$$

Eq.(13) shows the impact of gamma-ray energy E_γ , and formation atomic number Z on the mass attenuation coefficient μ_m , highlighting the response sensitivity among these variables. This work emphasizes the variability of the mass attenuation coefficient, which is different to previous studies where the mass attenuation coefficient was typically assumed to be constant. While changes in the mass attenuation coefficient within the average energy range of inelastic gamma rays in formations are small enough to have the effect of gamma energy ignored [37]. Instead, its close correlation with the formation's composition is emphasized, particularly the effects of lithology and pore content on the macroscopic atomic number(Z). To further prove this point, Fig.2 is shown which represents the difference in formation's macroscopic atomic numbers when the lithology and porosity content change. These changes in lithology and pore content indicate variations in the constituent elements of the formation, which directly affect the formation macroscopic atomic number(Z), resulting in alterations to the total mass attenuation coefficient. This affects gamma rays' attenuation, thereby complicating gamma-ray transport. So, the mass attenuation coefficient is treated as a function related to the formation lithology(l) and pore content(p). Accurate measurement of density can be achieved by describing the influence of different formation components on the gamma transport process.

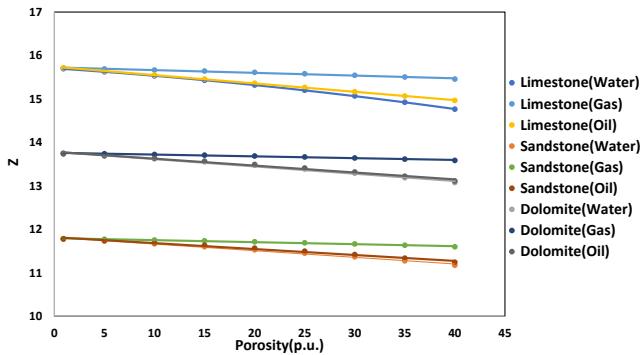


Fig. 2. (Color online) Z -results across different formations with various pore contents

According to the above analysis, treating mass attenuation coefficient as a function pertaining to environmental parameters will better depict gamma-formation reaction sensitivity and thus enable more accurate NGD calculations:

$$\mu_m(l, p) = c + d(Z(l, p) + 1) \quad (14)$$

$$c = \pi(r_0)^2 N_A \left\{ \frac{1 + \eta}{\eta^2} \left[\frac{2(1 + \eta)}{1 + 2\eta} - \frac{1}{\eta} \ln(1 + 2\eta) \right] + \frac{1}{2\eta} \ln(1 + 2\eta) - \frac{1 + 3\eta}{(1 + 2\eta)^2} \right\} \quad (15)$$

$$d = \frac{(r_0)^2 N_A}{274} \left(\frac{28}{9} \ln(2\eta) - \frac{218}{27} \right) \quad (16)$$

where c and d are constants.

After substituting the mass attenuation coefficient μ_m from Eq.(14) into Eq.(8), the equation can be reformulated as follows:

$$\rho = \frac{e \ln(RIN) + f \frac{1}{L_n}}{c + d(Z(l, p) + 1)} \quad (17)$$

This expression is composed of 2 key parameters: the fast neutron deceleration length L_n and the formation's macroscopic atomic number Z , and L_n is related to the formation density and the hydrogen index. These physical parameters cannot be directly measured, instead, they can be derived through the analysis of detector responses. Consequently, in the next section, we present a real pulse neutron logging tool and construct a high-fidelity Monte Carlo model for the analysis of these physical parameters.

C. Analysis of physical parameters using NGD tool

Geant4 (Geometry and Tracking 4), an open-source Monte Carlo platform, is used for simulations. The tool model, shown in Fig.3, has a total length of 2328 mm and a diameter of 188 mm, featuring four neutron detectors and two gamma detectors. To minimize the impact of water in the mud pipe on neutron detection, a boron-containing shield is positioned at the base of the neutron detectors. Additionally, the near gamma detector is used not only for density measurement but also for formation sigma and elemental measurements. To reduce interference from capture gamma rays generated by the tool's interaction with thermal neutrons, a two-layer shielding structure is implemented. A cubic space measuring $6 * 6 * 6$ m is designated to simulate formation environment, with a borehole diameter of 215.9 mm, positioning the tool at the center of the borehole. This tool is currently undergoing construction and will be deployed in the field upon completion. Therefore, it is selected to verify the feasibility of the proposed method. Extensive simulations are conducted using a NGD tool model, incorporating various formation lithologies (limestone, sandstone, dolomite) and porosity ranges (0-40 p.u.). These simulations aim to establish the relationship between detector responses and relevant physical parameters in the Eq. (17) and therefore will be used for concept validation.

The specific relationships are as follows:

(a) Hydrogen index(I_H)

The correlation between hydrogen index and detector responses is analyzed using simulation data. Fig.4(a) presents the correlation coefficients between various detector responses and hydrogen index. These coefficients measure the strength of the linear relationship between the variables, with values closer to 1 indicating a strong correlation. This analysis helps identify the optimal response for representing the hydrogen index. In Fig.4(a), the features represent various detector counts: $FN1$ and $FN2$ correspond to near and far fast neutron counts, $ETN1$ and $ETN2$ to near and far epithermal neutron counts, and $CAP1$ and $CAP2$ to near and far capture gamma counts. Additionally, RFN , $RETN$, and $RCAP$ represent the respective ratios of fast neutrons, epithermal neutrons, and capture gamma counts. As shown in

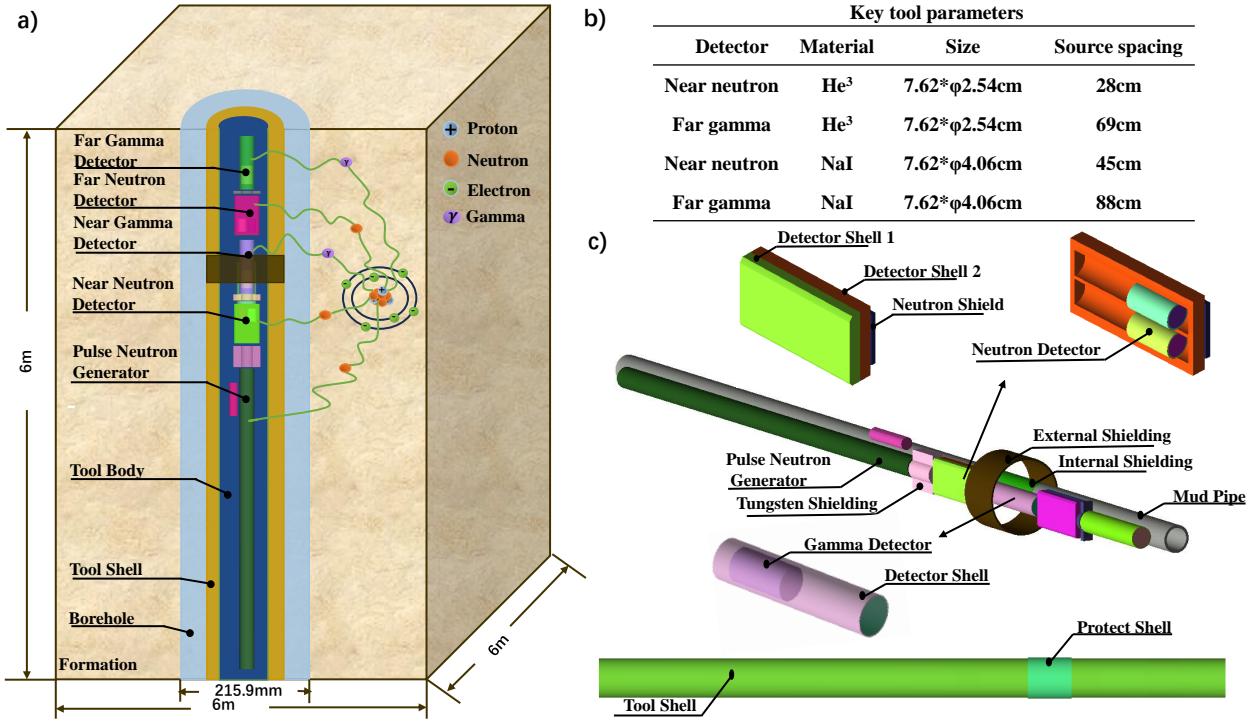


Fig. 3. (Color online) Tool model: (a) tool overview; (b) key tool parameters; (c) shielding structure

276 the figure, the ratio of near to far epithermal neutron counts
 277 ($RETN = ETN1/ETN2$) exhibit the strongest correlation
 278 with the hydrogen index, making $RETN$ the most effective
 279 indicator of hydrogen content among all detector responses.

280 (b) Formation density(ρ)

281 To accurately represent formation density, correlation anal-
 282 ysis is applied to evaluate the relationships between various
 283 detector responses and density. In Fig. 4(b), $IN1$ and $IN2$
 284 represent near and far inelastic gamma counts, while $CAP1$
 285 and $CAP2$ represent near and far capture gamma counts.
 286 RIN is the ratio of near to far inelastic gamma counts, ex-
 287 pressed as $RIN = IN1/IN2$. And $RCAP$ is the ratio of
 288 near to the far capture gamma counts. As shown in Fig.4(b),
 289 RIN exhibits the strongest correlation with density, making
 290 it the optimal parameter for describing density in Eq. (9).
 291 This is also consistent with NGD physics principle [38, 39].

292 (c) Formation macroscopic atomic number(Z)

293 The macroscopic atomic number Z , an inherent character-
 294 istic of the formation, is closely influenced by lithology and
 295 pore contents. By analyzing the detector counts within a spe-
 296 cific energy window (0.07 to 0.35 MeV), denoted as N_{lith} ,
 297 a relationship can be established to represent macroscopic
 298 atomic number. As shown in Fig. 4(c), this method allows for
 299 the derivation of Z from the detector responses, using counts
 300 within the designated energy range to effectively characterize
 301 the formation's macroscopic atomic number.

302 To summarize, density, hydrogen index and formation
 303 macroscopic atomic number Z can be represented using
 304 RIN , $RETN$ and N_{lith} , all of which can be obtained from
 305 detector counts. Since the root term in the represented equa-
 306 tion complicates the acquisition of calibration coefficients, a

307 polynomial fit approach was used to simplify the equation, as
 308 shown in Fig.4(d). After substituting the root term with a
 309 second-degree expression, previous Eq.(9) can be rewritten
 310 as:

$$311 \quad L_n = \frac{\tau_0}{\tau_1 \ln(RIN) * \sum_0^2 t_m (\ln(RETN))^m} \quad (18)$$

312 By substituting the L_n expression into Eq. (17), density
 313 equation can be obtained:

$$314 \quad \rho = \frac{\ln(RIN) * \sum_0^2 \beta_m (\ln(RETN))^m}{\theta_2 (N_{lith})^2 + \theta_1 N_{lith} + \theta_0} \quad (19)$$

315 where β_0 , β_1 , β_2 , θ_0 , θ_1 , θ_2 is the fitting parameter. From
 316 Eq. (19), the formation density is determined by three pa-
 317 rameters: the ratio of inelastic gamma counts RIN , the ratio
 318 of epithermal neutron counts $RETN$, and the count N_{lith} .
 319 The coefficients in the above equation using the Levenberg-
 320 Marquardt method are obtained.

III. RESULTS AND DISCUSSIONS

322 To prove the effectiveness of developing mass attenuation co-
 323 efficient function, we compare two approaches for treating
 324 the mass attenuation coefficient: as a constant versus as a
 325 function of formation composition. The results demonstrate
 326 that treating it as a function can significantly enhance calcula-
 327 tion accuracy, demonstrating the effectiveness of the method.
 328 Next, we assess the method's performance across various en-
 329 vironments, focusing on two critical factors: formation lithol-
 330

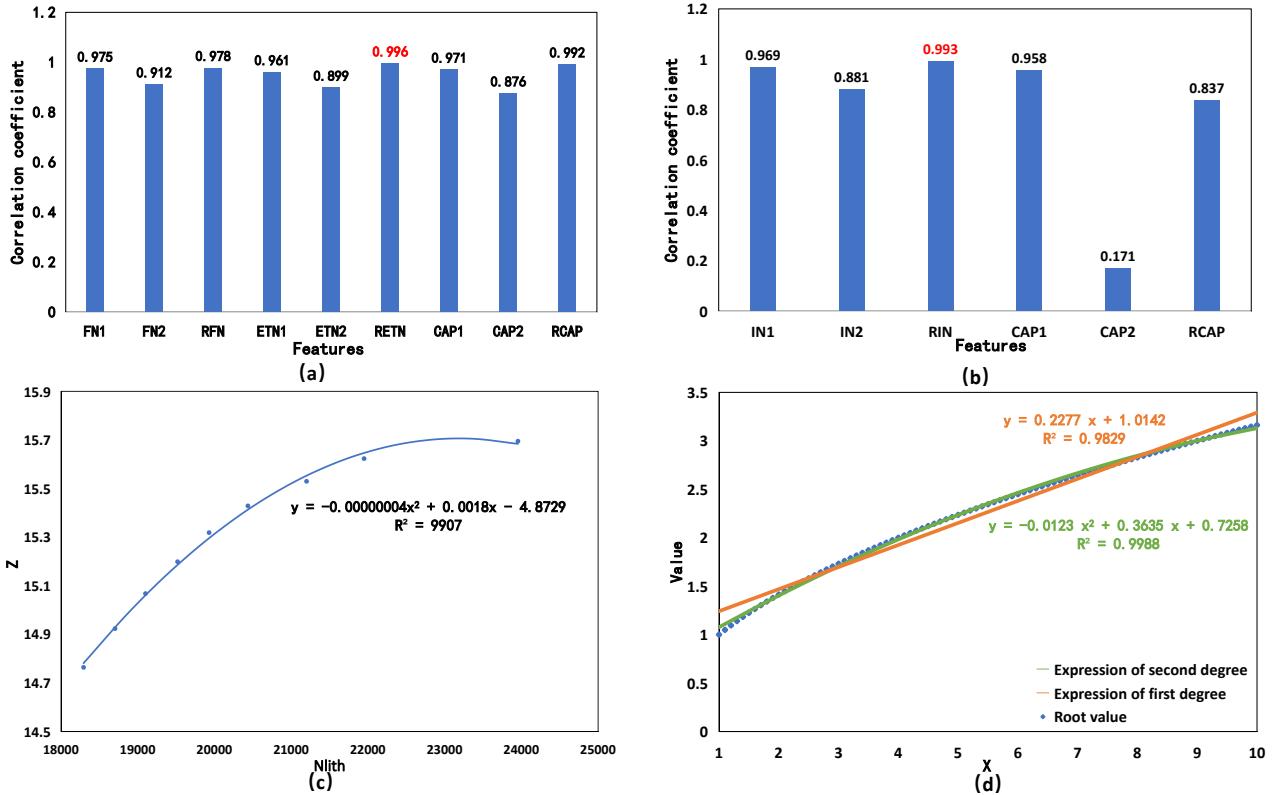


Fig. 4. (Color online) Analysis of physical parameters: (a) Hydrogen Index; (b) Density; (c) Limestone's Macroscopic Atomic Number; (d) Polynomial fit

ogy and pore content. Finally, we present three test cases to validate the method's applicability in complex formations. The absolute errors are used to evaluate the calculated density results, as expressed in Eq. (19). When the absolute errors are less than the threshold of 0.025 g/cm^3 , the calculated results are considered accurate [40].

$$337 \quad \text{Error} = |DT - DC| \quad (20)$$

338 where Error is the absolute error between calculated density(DC) and true density of the simulated 339 formation(DT).

341 A. Comparison of two approaches regarding mass 342 attenuation coefficient

343 Section 2.2 emphasizes that the proposed method treats the 344 mass attenuation coefficient as a function pertaining to for- 345 mation lithology and pore content. To evaluate the effective- 346 ness of this method, a comparison is conducted in this section, 347 primarily focusing on two approaches of the mass attenu- 348 ation coefficient: treating it as a constant (denoted as h) versus 349 treating it as a function. Based on Eq. (8) and the analysis 350 of the relevant physical parameters in Section 2.3, if the mass 351 attenuation coefficient is treated as h , the equation can be ob- 352 tained:

$$353 \quad \rho = \frac{\ln(RIN) * \sum_0^2 \beta_m (\ln(RETN))^m}{h} \quad (21)$$

354 where $\beta_0, \beta_1, \beta_2$ and h are the fitting parameters. From Eq. 355 (19), the density is determined by two parameters: the ratio 356 of inelastic gamma counts RIN and the ratio of epithermal 357 neutron counts $RETN$.

358 Limestone with densities ranging from 2.018 g/cm^3 to 359 2.862 g/cm^3 are designed to compare two approaches. Fig. 360 5(a) presents the absolute density errors of both approaches. 361 The results clearly illustrate significant differences: the con- 362 stant method has a relatively high average absolute error of 363 0.048 g/cm^3 , whereas the error calculated by the proposed 364 method is reduced by about four times compared to the con- 365 stant method, with an average absolute error of 0.012 g/cm^3 . 366 This demonstrates the effectiveness and accuracy of the new 367 method in measuring formation density.

368 B. Pore content impact analysis

369 In neutron gamma density(NGD) measurements, neutron 370 transport is sensitive to the presence of pore content. To eval- 371 uate the impact of different pore contents on the accuracy of 372 this method, limestone with porosities ranging from 0.9 to 40 373 p.u. are selected, with pores filled with water, gas, or oil. The 374 density results under different pore contents are shown in Ta-

ble 1 and Fig.5 (b).

TABLE 1. Density results across different pore fluids

Porosity (p.u.)	Water(1.0 g/cm ³) Error (g/cm ³)	Gas(0.2 g/cm ³) Error (g/cm ³)	Oil(0.835 g/cm ³) Error (g/cm ³)
0.9	0.010	0.003	0.005
10	0.005	0.007	0.019
20	0.002	0.003	0.007
25	0.003	0.012	0.020
30	0.006	0.011	0.010
35	0.008	0.011	0.012
40	0.008	0.005	0.012

Table 1 and Fig.5(b) present the density calculation results for varying pore contents. For analysis, seven porosity types are selected, each tested under conditions where the pores are filled with water, oil, and gas. In limestone with pores filled with water, the Hydrogen Index I_H is equivalent to the formation's porosity. The results indicate that the I_H has a minimal impact on density measurements. Regardless of I_H variations, the errors between calculated densities and true densities are less than the threshold of 0.025 g/cm³, demonstrating that densities calculated by the new method consistently align well with true densities. Additionally, the method can also achieve accurate measurements in high I_H formations. When comparing density calculations for different pore contents, it is observed that when the pores are filled with water or gas, the absolute density errors are relatively small, remaining below 0.015 g/cm³. However, the errors are relatively large when the pores are filled with oil. Notably, whether the pores are filled with water, oil, or gas, the absolute errors are less than 0.02 g/cm³. This demonstrates that the method can accurately calculate formation density across various porosities and pore contents.

Since different lithologies affect neutron transport and gamma attenuation, 42 models, including limestone, sandstone, dolomite, and one-to-one mixture of any two lithologies, are designed to verify the accuracy of the proposed method. All model pores are filled with water, with porosities ranging from 0.9 to 40 p.u. The densities calculated using the proposed method are compared with true densities used in simulated model construction, which varied between 1.93 g/cm³ and 2.843 g/cm³. The density results are shown in Fig.5(c) and Table 2.

To verify the impact of lithology on density measurement, the study focuses on two types of formations: single lithology (clean formations) and mixed lithology composed of sandstone, limestone, and dolomite. As shown in Fig.5(c) and Table 2, there are slight differences in density results across various lithologies, and the average absolute error in mixed lithology is slightly larger than that in single lithology, likely due to the complexity of the formation's composition. Whether it is a single lithology or a mixed lithology, the calculated densities closely align with the true densities, and the absolute

density errors less than 0.02 g/cm³. Overall, the average absolute error in the test database is 0.009 g/cm³, confirming the accuracy of the proposed method under different lithologies.

D. Multi-parameter impact analysis

The above results quantitatively analyze the impact of lithology and pore content on the accuracy of the density measurement. To further evaluate the proposed method, three cases (Case 1, Case 2, and Case 3) are designed, representing three lithologies (limestone, sandstone, dolomite) and three pore contents (water, oil, gas), while considering mud components (such as chlorite). Specifically, Case 1 simulates a water-filled limestone with 20% chlorite, Case 2 simulates an oil-filled sandstone with 10% chlorite, and Case 3 simulates dolomite containing gas, which has a relatively high chlorite content of 40%. In all cases, the borehole size is 8.5 inches, with a porosity range set from 0 to 25 p.u., and formation densities between 2.211 g/cm³ and 2.712 g/cm³. The results are illustrated in Fig.6.

From Fig.6, the inelastic gamma ratio and epithermal neutron ratio vary across formations with different lithologies and pore contents, highlighting the impact of formation composition on neutron transport and gamma attenuation. In particular, Case 1 and Case 2 exhibit smaller measurement errors, demonstrating higher accuracy. In contrast, Case 3 exhibits relatively higher measurement errors, possibly due to its more complex formation composition characterized by elevated mud content and gas-filled pores. Nevertheless, the absolute density errors in all three cases remain below 0.02 g/cm³, demonstrating the accuracy and reliability of the proposed method for measuring formation density.

IV. CONCLUSIONS

C. Lithology impact analysis

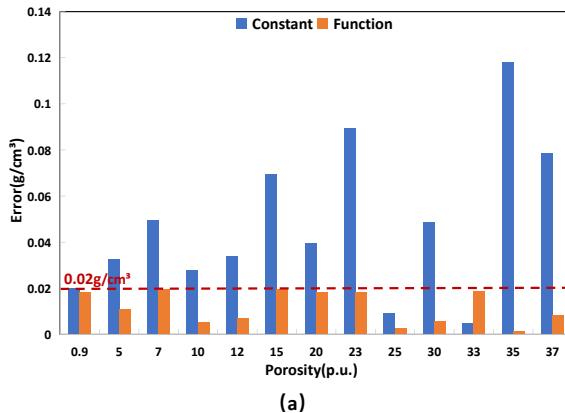
(1) A new mass attenuation coefficient function of formation lithology and pore content is introduced. Based on neutron induced gamma attenuation process study, mass attenuation coefficient is shown varying pertaining to formation parameters, therefore, this work proposes to consider it as a function to better evaluate the effects of environmental variables regarding gamma attenuation.

(2) A new density measurement method is developed by employing the concept of mass attenuation coefficient function that evaluates the effects of formation composition on gamma attenuation. The method relies on inelastic gamma rays for density measurement while incorporating epithermal neutrons to correct neutron transport, for example, fast neutron influences on spatial distribution and intensity of inelastic gamma rays. By integrating information from both neutrons and gamma rays, this method can evaluate interaction mechanisms between radiation and formation more accurately and therefore helps obtain more precise density measurement.

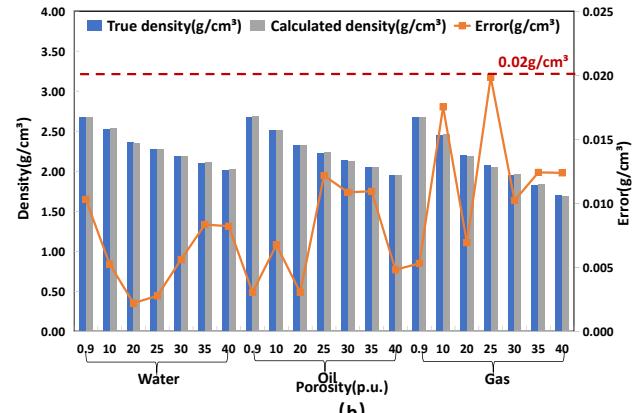
TABLE 2. Density results across different lithologies

Porosity (p.u.)	Limestone(g/cm ³)		Sandstone(g/cm ³)		Dolomite(g/cm ³)	
	True Density	Error	True Density	Error	True Density	Error
0.9	2.682	0.010	2.682	0.011	2.843	0
10	2.527	0.005	2.485	0.011	2.767	0.001
20	2.358	0.002	2.320	0.011	2.674	0.001
25	2.273	0.003	2.238	0.009	2.488	0.002
30	2.188	0.006	2.155	0.007	2.395	0.001
35	2.103	0.008	2.073	0.005	2.209	0.010
40	2.018	0.008	1.990	0.009	2.116	0.003

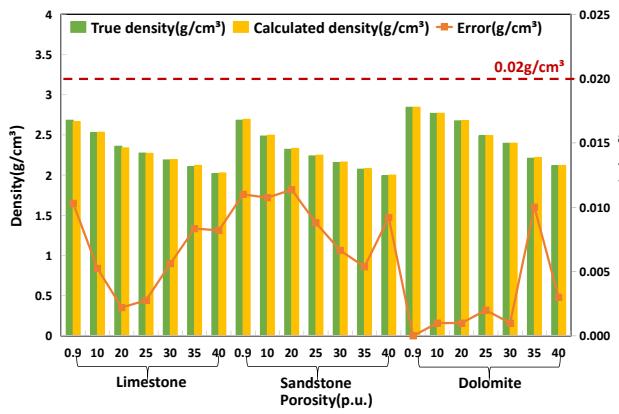
Porosity (p.u.)	Limestone+Sandstone(g/cm ³)		Limestone+Dolomite(g/cm ³)		Sandstone+Dolomite(g/cm ³)	
	True Density	Error	True Density	Error	True Density	Error
0.9	2.658	0.011	2.763	0.019	2.739	0.007
10	2.506	0.011	2.601	0.016	2.580	0.004
20	2.339	0.019	2.423	0.004	2.404	0.007
25	2.255	0.005	2.334	0.005	2.316	0.014
30	2.172	0.003	2.245	0.011	2.229	0.020
35	2.088	0.005	2.156	0.015	2.141	0.006
40	2.004	0.014	2.067	0.006	2.053	0.007



(a)



(b)



(c)

Fig. 5. (Color online) Density results: (a) Two approaches comparison; (b) Pore content results; (c) Lithology results

(3) An elaborate NGD tool model is built and employed to verify the performance of the new method. The proposed method is evaluated using a total of 63 sets of simulated models of varying lithologies and pore contents. The results show that the absolute errors of density calculated by the method are below 0.02 g/cm³ for

all cases. Specifically, the method obtains the same level of accuracy in mixed cases, proving its effectiveness. This can offer theoretical support for the design of new NGD tools. The method faces challenges in terms of extreme environmental conditions and tool calibration. For example, under logging-while-drilling

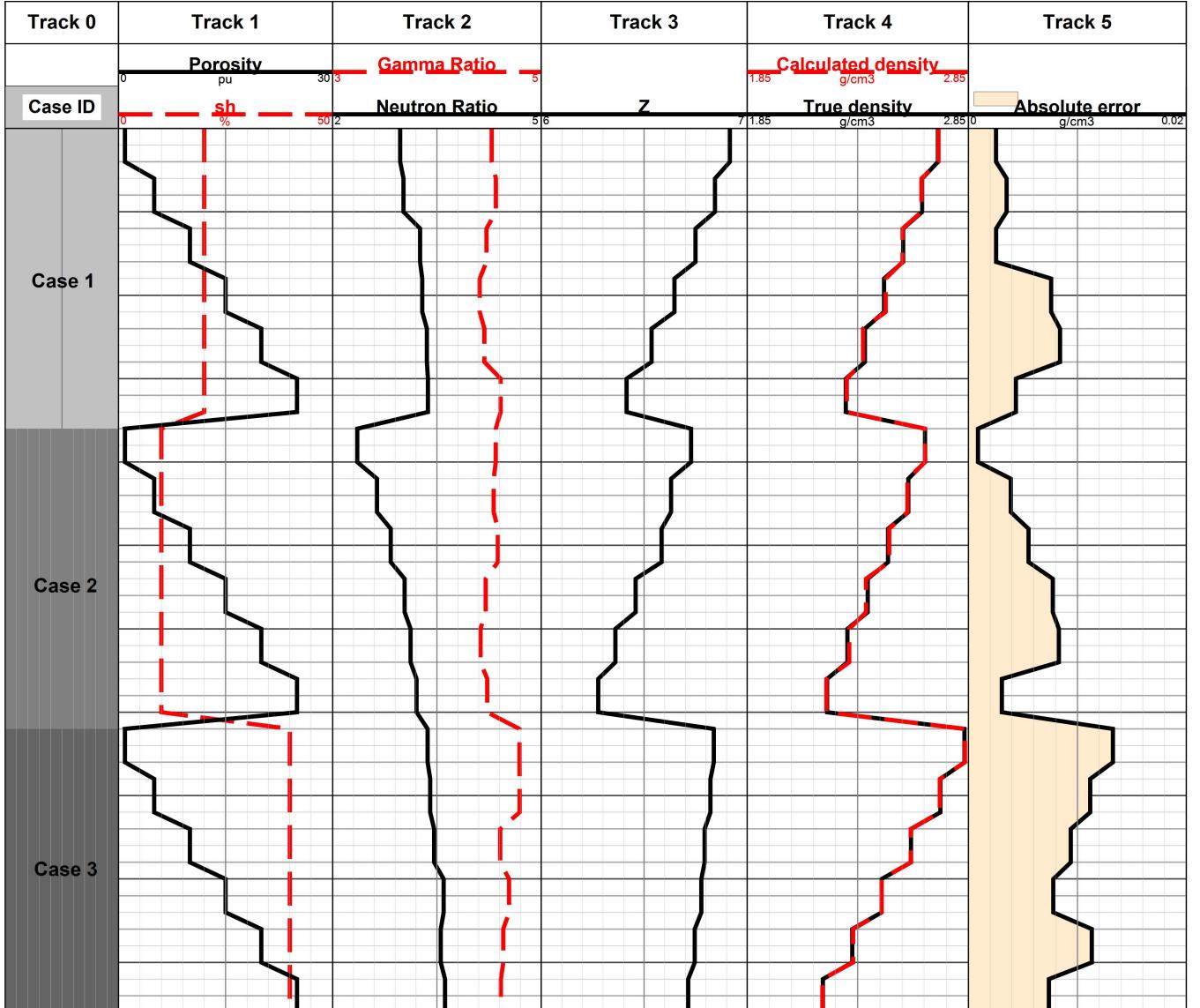


Fig. 6. (Color online) Results and comparisons. Track 0: Three cases, of which Case 1 is limestone filled with water, Case 2 is sandstone filled with oil, and Case 3 is dolomite filled with gas. Track 1: Case parameters, including porosity and mud content in the formation. Track 2: Detector counts, including the inelastic gamma count ratio $LNRIN$, and the epithermal neutron count ratio $LNRETN$. Track 3: Formation's macroscopic atomic number. Track 4: Comparison of the true density with the calculated density. Track 5: Absolute error.

487 downhole conditions such as high temperature (150°C) 497
 488 and high pressure (2000 psi), the performance of the
 489 tool's detectors and electronics may be affected, while
 490 changes in the physical properties of borehole fluids
 491 can also occur, thus impacting measurement accuracy.
 492 Additionally, the method relies on significant amounts
 493 of calibration coefficients, this requires high-level cali-
 494 bration standards in tool-specific environments. To im-
 495 prove the applicability of the method, further in-depth
 496 research will be conducted regarding these two aspects.

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NOMENCLATURE

501 μ_m	Total mass attenuation coefficient
502 μ_{co}	Mass attenuation coefficient for Compton effect
503 μ_{pa}	Mass attenuation coefficient for Pair production
504 ϕ_n	Fast neutron distribution

505	Φ_{in}	The inelastic gamma-rays distribution	525	$IN1$	Near inelastic gamma count
506	ρ	Formation density	526	$IN2$	Far inelastic gamma count
507	Σ_{in}	Inelastic scattering cross-section	527	l	Formation lithology
508	A	The atomic weight	528	L_n	Fast neutron deceleration length
509	c	The speed of light in a vacuum	529	L_1	The distance of near gamma detector
510	$CAP1$	Near captured gamma count	530	L_2	The distance of far gamma detector
511	$CAP2$	Far captured gamma count	531	m_e	The electron rest mass
512	D_n	Neutron diffusion coefficient	532	N_A	Avogadro's constant
513	DC	Calculated density	533	NGD	Neutron-gamma density
514	DT	True density of the simulated formation	534	p	Pore content
515	E_0	Initial energy of fast neutron	535	Q	The number of neutrons emitted per second
516	E_γ	Gamma-ray energy	536	r	The distanc between source and detector
517	$ETN1$	Near epithermal neutron count	537	r_0	The classical electron radius
518	$ETN2$	Far epithermal neutron count	538	$RCAP$	Ratio of near to far capture gamma counts
519	$FN1$	Near fast neutron count	539	$RETN$	Ratio of near to far epithermal neutron counts
520	$FN2$	Far fast neutron count	540	RFN	Ratio of near to far fast neutron counts
521	GGD	Gamma-gamma density	541	RIN	Ratio of the near to far inelastic gamma counts
522	i	The average number of inelastic gamma rays after neutron enters the formation	542	$Sigma$	Formation macroscopic capture cross section
523			543	Z	Atomic number
524	I_H	Hydrogen index	544		

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